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Sample Collection			
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<b>Patient Data</b> Name					5.2.0.13
Name					
	MRS. MANISHA		Patient ID		012111270137
Birthday		15/05/1999	Sample ID		11169714
Age at delivery		23.08	Sample Date		27/11/2021
Gestational age		12+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	57 Diabetes		NO	Pregnancies	unknown
Smoker N	VO Origin		Asian		
Biochemical Data		Ultrasound Da	ata		
Parameter Val	ue (	Corr Mom	Gestational age	2	11+6
PAPP-A 1.	86 mIU/ml	0.46	Method		CRL(<>Robinson
fb-hCG 23	3.5 ng/ml	0.51	Scan date		27/11/2021
Risks at sampling date			Nuchal translucency 1.2		
Age Risk	]	1:1462	Nuchal translu	cency MoM	0.83
Biochemical T21 risk	]	1:5505	Nasal bone		Present
Combined T21 risk		<1:10000			
Trisomy 13/18+NT		<1:10000			
Risk		Down's Syndrome Risk (Trisomy 21 Screening)			
Risk   1:10   1:250   Cut off   1:1000   1:1000   1:1000   1:1000   1:1000   1:1000   1:1000   1:1000   1:1000   1:1000   1:10000   1:10000, which indicates a low risk			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value! The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values Risk above Age Risk Risk below Age risk		